

SPECIFICATION

Drug and food or drink for improving hyperglycemia

Technical Field

The present invention relates to a drug and food or drink for improving hyperglycemia, which contains a compound that can be safely ingested without causing acute hypoglycemia and has a long-term blood glucose level control action lowering the hemoglobin A1c level.

Background Art

Hemoglobin A1c, a binding product of glucose and hemoglobin, increases depending on the severity of hyperglycemia in a glucose level-dependent manner. Because hemoglobin A1c once produced is not eliminated until the lifetime of erythrocyte (120 days) runs out, it reflects the past blood glucose control conditions over a long period of time (Non-patent document 1). Hemoglobin A1c was adopted as a selected test item of the basic health screening according to the Health Law for the Aged since 1996 and adopted as an auxiliary diagnosis indicator of diabetes mellitus in the new diagnosis criteria of diabetes mellitus in 1999. Therefore, it is considered that hemoglobin A1c is an indicator of great clinical significance (Non-patent document 2).

If a hyperglycemic condition is sustained, glucose specific insulin hyposecretion and insulin resistance are observed and serve as factors that further aggravate hyperglycemia (Non-patent document 3). Because long-term blood glucose level control is necessary to prevent progression from the hyperglycemic condition to onset of

diabetes mellitus, it is considered to become necessary to suppress increase in the hemoglobin A1c level.

Alimentotherapies and exercise are recommended to control blood glucose level in patients with prediabetes (condition suspected of diabetes). Although various functional foods for preventing postprandial increases in blood glucose level (food for specified health uses) have already been marketed, all of these only have a temporary effect of suppressing increase in blood glucose level. Therefore, control of blood glucose level over a long period of time cannot be expected, and development of such a substance having a hemoglobin A1c level lowering action has been desired.

Furthermore, α -glucosidase inhibitors, sulfonylurea drugs as insulin secretagogues, thiazolidine derivatives as insulin resistance improving agents and so forth are currently used as therapeutic agents for diabetes mellitus. However, the drug efficacies thereof are not satisfactory, and they suffer many problems such as side effects causing coma due to rapid drop in blood glucose level.

Under the aforementioned circumstances, discovery of a substance that can be safely ingested without causing acute hypoglycemia and has a long-term blood glucose level control action by decreasing the hemoglobin A1c level has been strongly desired.

Conventionally, as examples of substances having an effect of suppressing increases in blood glucose level, the prior art references have disclosed a hyperglycemia suppressing agent containing a banaba-derived ingredient (Patent document 1), a hyperglycemia suppressing agent containing a concentrated extract of fermentation product of wheats or barleys as an active ingredient (Patent

document 2) and so forth.

Furthermore, as techniques of using a triterpene glycoside as an active ingredient, for example, a diabetes preventing agent containing a glycoside extracted from *Gymnema inodorum* as an active ingredient (Patent document 3), a metabolism improving method and a composition therefor containing corosolic acid extracted from banaba as an active ingredient (Patent document 4), a lipase inhibitor (Patent document 5) and a triterpene derivative having an immunosuppressing activity (Patent document 6) have been disclosed.

Furthermore, it has been disclosed that the insulin action enhancing activity of a compound having a lanostane skeleton or 3,4-secolanostane skeleton (Patent document 7) enhances the insulin action in regulation of adipocyte differentiation, although the effect thereof on diseases in the pancreas is unknown.

Furthermore, compounds selected from the group consisting of 24-alkylcholesten-3-ones and 24-alkylcholestan-3-ones that have no double bond in the basic steroid skeleton have been disclosed as hypoglycemic agents (Patent document 8).

Furthermore, as the prior arts concerning compounds having the cyclolanostane skeleton, a method for producing cyclobranol or cyclobranol ferulic acid ester (Patent document 9) as well as tranquilizers (Patent document 10), hypolipidemic drugs (Patent document 11), interferon inducers (Patent document 12), ovulation inducing agents (Patent document 13) and oncogenesis preventive drugs (Patent document 14) containing 24-methylenecycloartanol as an active ingredient have been disclosed. However, effects of compounds having the cyclolanostane skeleton on blood

glucose levels and hemoglobin Alc levels are not mentioned in these references.

The genus *Aloe* in the family *Liliaceae* is a group of plants including *Aloe vera* (*Aloe barbadensis* Miller) and *Aloe arborescens* (*Aloe arborescens* Miller var. *natalensis* Berger) and so forth, and they are empirically known to have various efficacies. The prior arts regarding the use of plants of the genus *Aloe* include immunomodulating polysaccharides (Patent document 15), immunosuppression improving agents containing a butanol fraction of an aloe extract or aloin (Patent document 16), HSP60 family protein synthesis suppressing agents containing aloin derivatives (Patent documents 17 to 19), protein having lectin activity derived from aloe leaf-skin (Patent document 20) and so forth.

As the prior arts regarding improvement of blood glucose levels by the plants of the genus *Aloe*, clinical studies in the United States (Non-patent document 4) and a hypoglycemic action observed in animal studies (Non-patent documents 5 and 6 and polysaccharides in plants of the genus *Aloe* (Patent document 21) have been disclosed. In these prior arts, the hypoglycemic ingredients of the plants of the genus *Aloe* were predicted to be polysaccharides or glycoproteins. Furthermore, it has been disclosed that, in a pressed extract of *Aloe vera* and a hypoglycemic agent containing the extract as an active ingredient (Patent document 22), a characteristic peak unique to an ester group observed in the FT-IR chart correlates with the activity, that the active ingredient is a polysaccharide, amino acid, malic acid or the like, and that the aforementioned active ingredient is degraded in commercially available *Aloe vera* gel powders, *Aloe vera* gel

solutions and Aloe vera gel extracts. Further, in addition to the above, a hypoglycemic action of aloe polysaccharides (Patent document 23), antioxidative action of 7-hydroxychromone contained in aloe (Patent document 24), a method for producing cycloartanol from cacao shells (Patent document 25) and so forth have been disclosed.

[Patent document 1] Japanese Patent Laid-open (Kokai) No. 2003-095941

[Patent document 2] Japanese Patent Laid-open No. 2002-371003

[Patent document 3] Japanese Patent Laid-open No. 05-247086

[Patent document 4] Japanese Patent Laid-open No. 2002-205949

[Patent document 5] Japanese Patent Laid-open No. 09-040689

[Patent document 6] International Patent Unexamined Publication in Japanese (Kohyo) No. 11-511482

[Patent document 7] Japanese Patent Laid-open No. 10-330266

[Patent document 8] Japanese Patent Laid-open No. 2003-048837

[Patent document 9] Japanese Patent Laid-open No. 50-160262

[Patent document 10] Japanese Patent Laid-open No. 55-153719

[Patent document 11] Japanese Patent Laid-open No. 59-027824

[Patent document 12] Japanese Patent Laid-open No. 59-036623

[Patent document 13] Japanese Patent Laid-open No. 59-073600

[Patent document 14] Japanese Patent Laid-open No. 2003-277269

[Patent document 15] International Patent Application Unexamined Publication in Japanese No. 2001-520019

[Patent document 16] Japanese Patent Laid-open No. 08-208495

[Patent document 17] Japanese Patent Laid-open No. 10-120576

[Patent document 18] Japanese Patent Laid-open No. 10-045604

[Patent document 19] Japanese Patent Laid-open No. 10-036271

[Patent document 20] Japanese Patent Laid-open No. 09-059298

[Patent document 21] Japanese Patent Laid-open No. 60-214741

[Patent document 22] Japanese Patent Laid-open No. 2003-286185

[Patent document 23] U.S. Patent No. 4,598,069

[Patent document 24] U.S. Patent Application Publication No. 2003/0207818

[Patent document 25] U.S. Patent Application Publication No. 2002/0048613

[Non-patent document 1] Nippon Rinsho, No. 748, Vol. 1, pp.615-617, 1999

[Non-patent document 2] Nippon Rinsho, No. 808, Vol. 2, pp.405-409, 2002

[Non-patent document 3] New England Journal of Medicine, Vol. 329, pp.977-986, 1993

[Non-patent document 4] Phytomedicine, Vol. 3, pp. 245-248, 1996

[Non-patent document 5] Phytotherapy Research, Vol. 15, pp.157-161, 2001

[Non-patent document 6] Phytotherapy Research, Vol. 7, pp.37-42, 1993

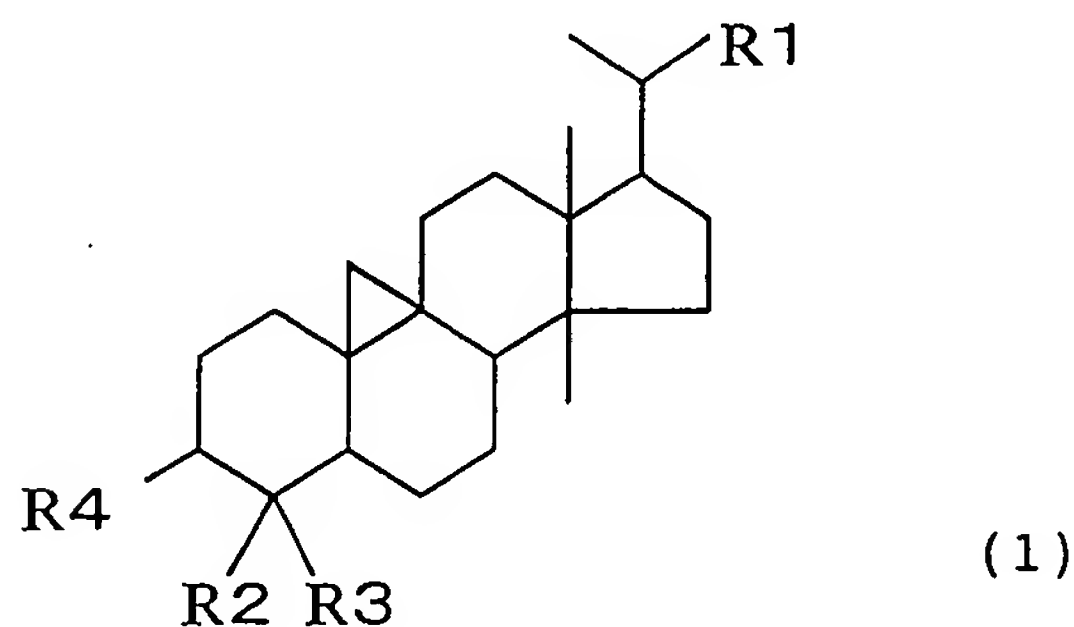
SUMMARY OF THE INVENTION

An object of the present invention is to provide a drug and food or drink for improving hyperglycemia, which comprises a compound that can be safely ingested without causing acute hypoglycemia and has a long-term blood glucose level control action lowering the hemoglobin A1c level.

The inventors of the present invention assiduously studied in order to achieve the foregoing object. As a result, they found that a compound having the cyclolanostane skeleton could be safely ingested without causing acute hypoglycemia and had a long-term blood glucose level control action lowering the hemoglobin A1c level. The present invention was accomplished on the basis of the above finding.

That is, the present invention provides a drug and food or drink for improving hyperglycemia, which comprises a compound having the cyclolanostane skeleton as an active ingredient.

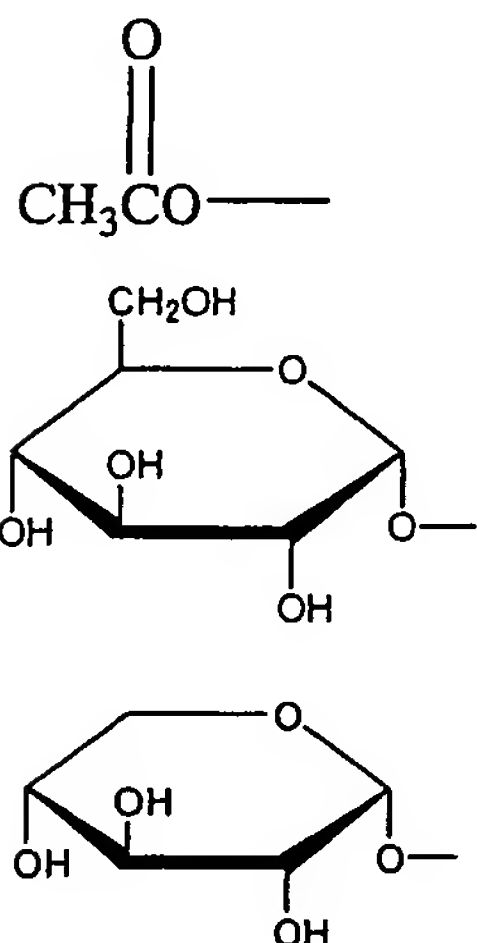
More specifically, the present invention provides a drug and food or drink for improving hyperglycemia, which comprises a compound represented by the following general formula (1) as an active ingredient.



In the formula, R1 represents a straight or branched

alkyl group having 6 to 8 carbon atoms, which may contain no double bond or 1 or 2 double bonds and may contain no hydroxyl group or carbonyl group or 1 or 2 hydroxyl groups and/or carbonyl groups, R2 and R3 each independently represent a hydrogen atom or a methyl group, and R4 forms C=O with the carbon atom constituting the ring or is a group represented by any one of the following formulas.

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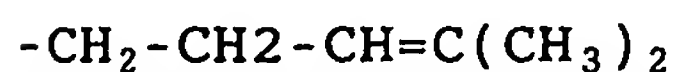
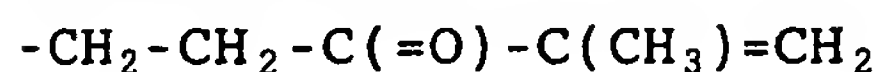
According to a preferred embodiment of the aforementioned drug and food or drink, R2 and R3 of the aforementioned compound both are methyl groups, and R4 is a hydroxyl group. Further, according to a preferred embodiment of the aforementioned drug and food or drink, R1 of the aforementioned compound is represented by any one of the following formulas.



(wherein Ra is any of hydrogen atom, hydroxyl or methyl group, and Rb is hydrogen atom or hydroxyl group)



(wherein R_c is any of hydrogen atom, hydroxyl or methyl group)



Further, according to a particularly preferred embodiment of the aforementioned drug and food or drink, the aforementioned compound is 9,19-cyclolanostan-3-ol or 24-methylene-9,19-cyclolanostan-3-ol.

Further, according to a preferred embodiment, the aforementioned drug contains 0.001 to 10% by dry mass of the aforementioned compound.

Further, according to a preferred embodiment, the aforementioned food or drink contains 0.0001 to 1% by dry mass of the aforementioned compound.

The present invention further provides a drug for improving hyperglycemia, which comprises an organic solvent extract or hot water extract of a plant or a fraction thereof as an active ingredient and contains 0.001 to 10% by dry mass of a compound represented by the aforementioned general formula (1) and, or food or drink for improving hyperglycemia, which comprises an organic solvent extract or hot water extract of a plant or a fraction thereof as an active ingredient and contains 0.0001 to 1% by dry mass of a compound represented by the aforementioned general formula (1). The aforementioned plant is preferably a plant of the family *Gramineae* or *Liliaceae*, and according

to a particularly preferred embodiment, the aforementioned plant of the family *Liliaceae* is a plant classified into the genus *Aloe*.

The present invention further provides the aforementioned food or drink attached with an indication that it is used for improvement of hyperglycemia.

Hereafter, the aforementioned drug and food or drink are also generically referred to as "the drug or food or drink of the present invention."

The present invention further provides use of a compound represented by the aforementioned general formula (1) or a composition containing the same for the production of a drug for improving hyperglycemia. According to a preferred embodiment of the use of the present invention, the aforementioned compound or composition containing the same contains 0.001 to 10% by dry mass or more of the aforementioned compound.

The present invention further provides a method for improving hyperglycemia, which comprises administering a compound represented by the aforementioned chemical formula (1) or a composition containing the same to a subject whose hyperglycemia is to be improved. According to a preferred embodiment of the method of the present invention, the aforementioned composition contains 0.001 to 10% by dry mass or more of the aforementioned compound.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a graph showing changes over time in random blood glucose levels during the repetitive administration period of samples. "o" denotes the results for the test sample 1-administered group, "Δ" denotes the results for

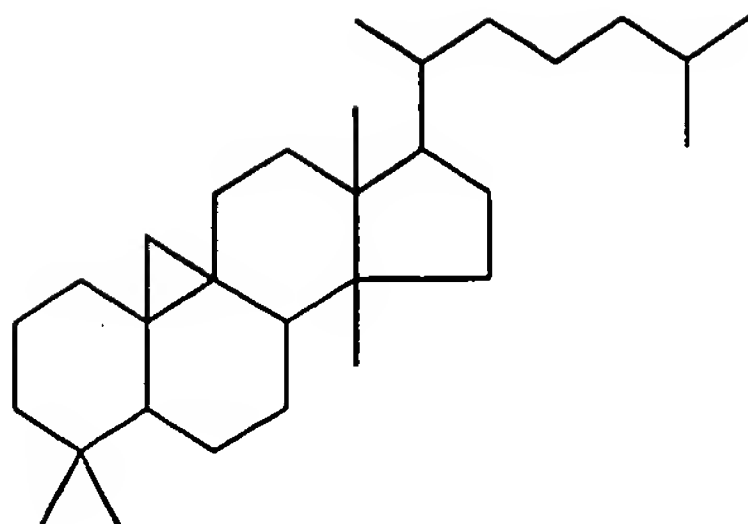
the test sample 2-administered group, "●" denotes the results for the negative sample-administered group, and "■" denotes the results for the control sample-administered group.

Fig. 2 is a graph showing changes over time in fasting blood glucose levels during the repetitive administration period of samples. "○" denotes the results for the test sample 1-administered group, "Δ" denotes the results for the test sample 2-administered group, "●" denotes the results for the negative sample-administered group, and "■" denotes the results for the control sample-administered group.

BEST MODE FOR CARRYING OUT THE INVENTION

Hereafter, preferred embodiments of the present invention will be explained in detail. However, the present invention is not limited to the following preferred embodiments, and the preferred embodiments can be freely modified within the scope of the present invention.

According to an embodiment, the drug or food or drink of the present invention contains a compound having the cyclolanostane skeleton and having a hyperglycemia improving effect and a hemoglobin Alc lowering action (hereinafter also referred to as "the compound of the present invention") as an active ingredient. The cyclolanostane skeleton refers to a compound represented by the following general formula (2).

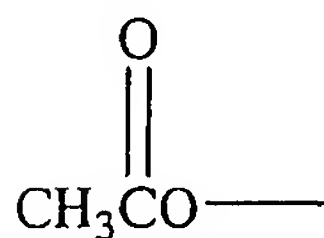


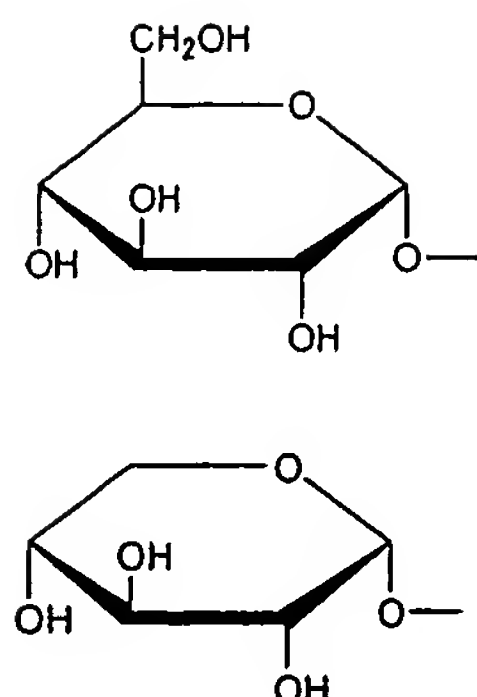
(2)

Specific examples of the compound having the cyclolanostane skeleton include compounds represented by the aforementioned general formula (1). The number of double bonds existing in the compound having the cyclolanostane skeleton is not particularly limited. Further, the number of double bonds existing in the ring is not particularly limited either. When 2 or more double bonds exist, they may be conjugated. The drug or food or drink of the present invention may contain 2 or more types of the compound of the present invention.

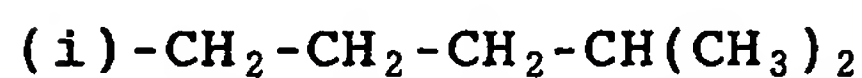
In the compound of the present invention of the aforementioned general formula (1), R1 represents a straight or branched alkyl group having 6 to 8 carbon atoms, which may contain no double bond or 1 or 2 double bonds and may contain no hydroxyl group or carbonyl group or 1 or 2 hydroxyl groups and/or carbonyl groups, R2 and R3 each independently represent a hydrogen atom or a methyl group, and R4 forms C=O with the carbon atom constituting the ring or is a group represented by any one of the following formulas.

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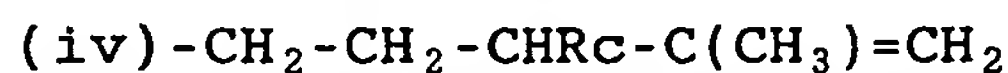
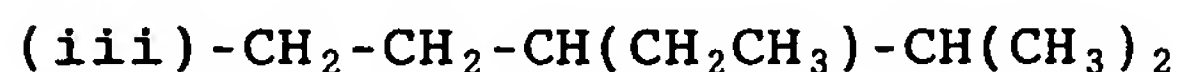




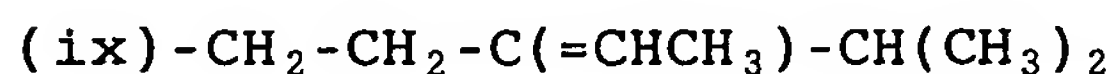
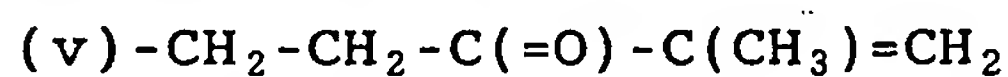
In the aforementioned general formula (1), R1 is preferably any one of the groups represented by the following formulas.



(wherein Ra is any of hydrogen atom, hydroxyl or methyl group, and Rb is hydrogen atom or hydroxyl group)



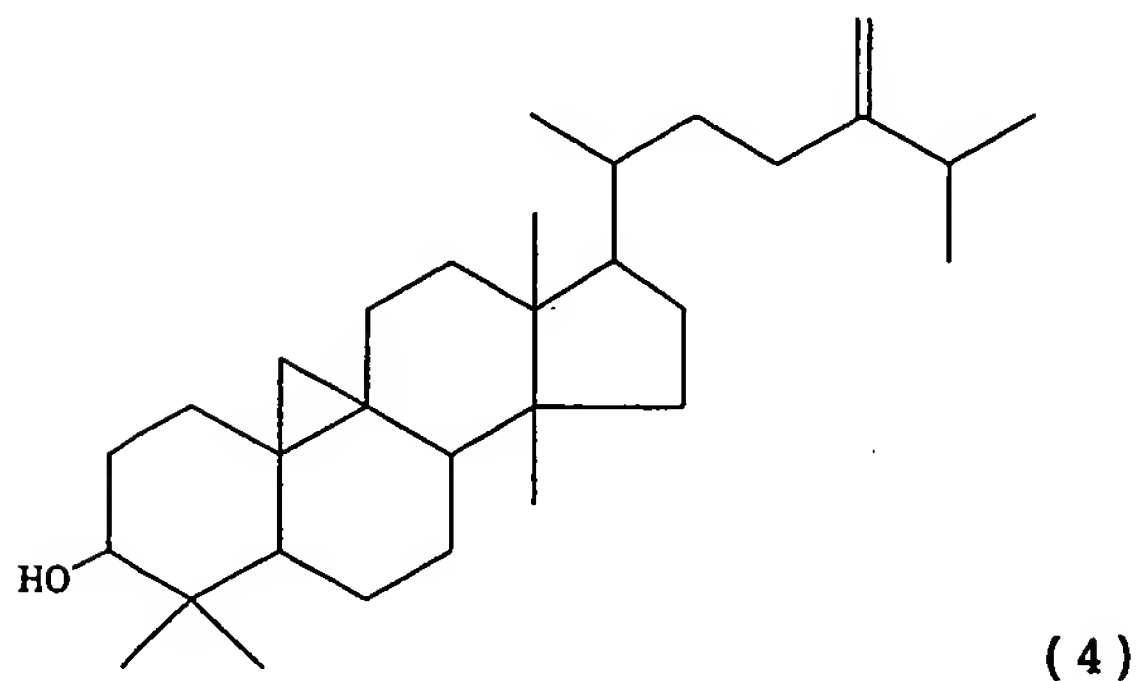
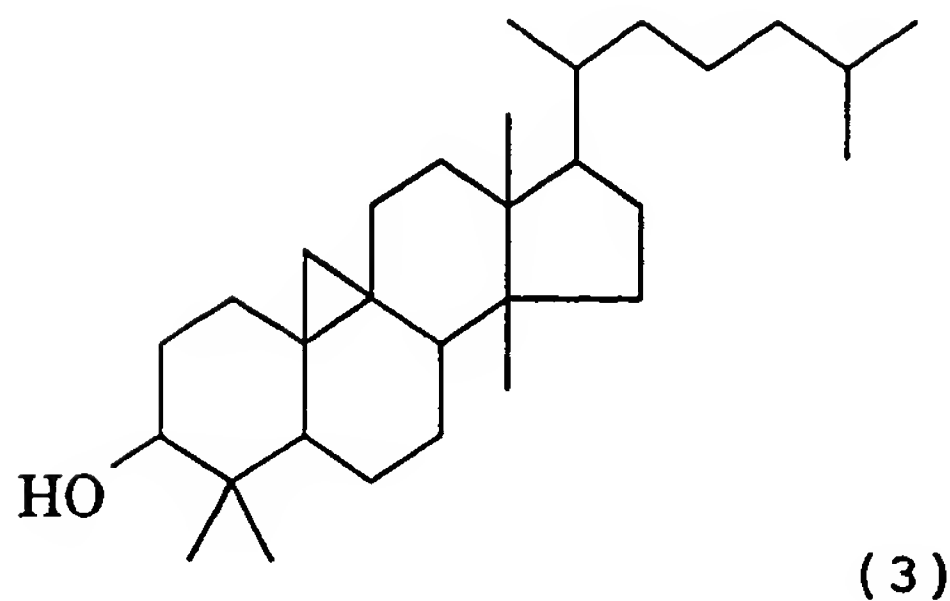
(wherein Rc is any of hydrogen atom, hydroxyl or methyl group)



Further, in the aforementioned general formula (1), it is preferred that R2 and R3 are both methyl groups, and R4 is a hydroxyl group.

The most preferred compounds as the aforementioned compound are those represented by the following formulas, 9,19-cyclolanostan-3-ol (formula (3)) and 24-methylene-

9,19-cyclolanostan-3-ol (formula (4)).

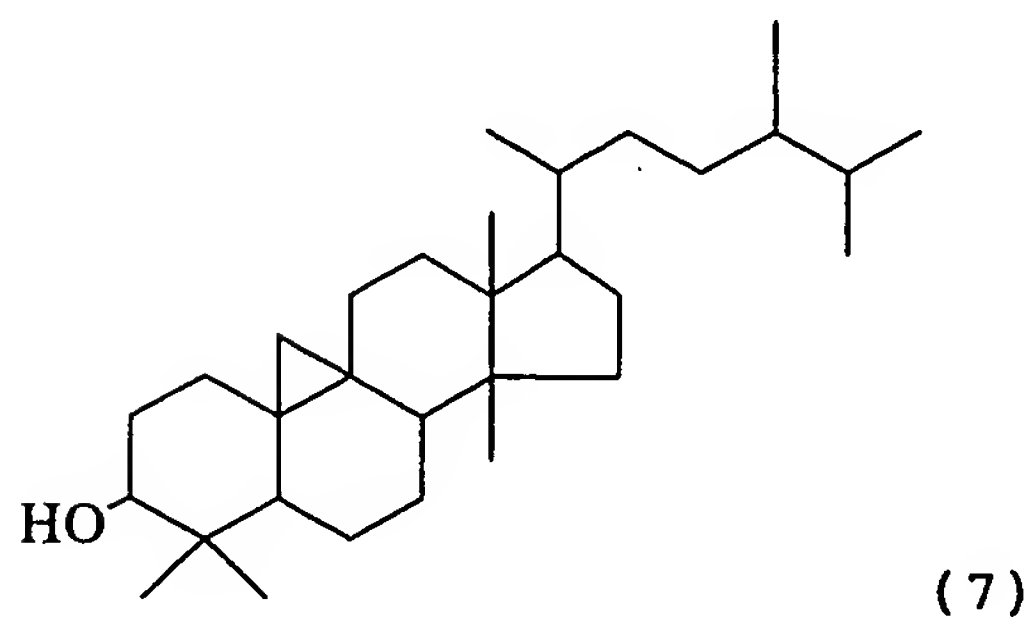
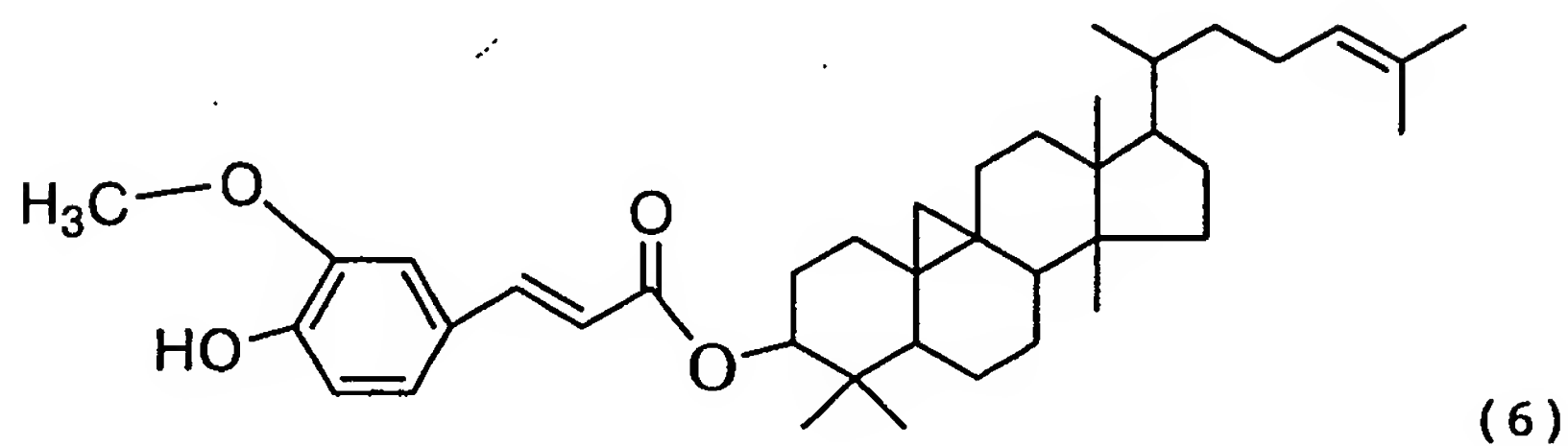
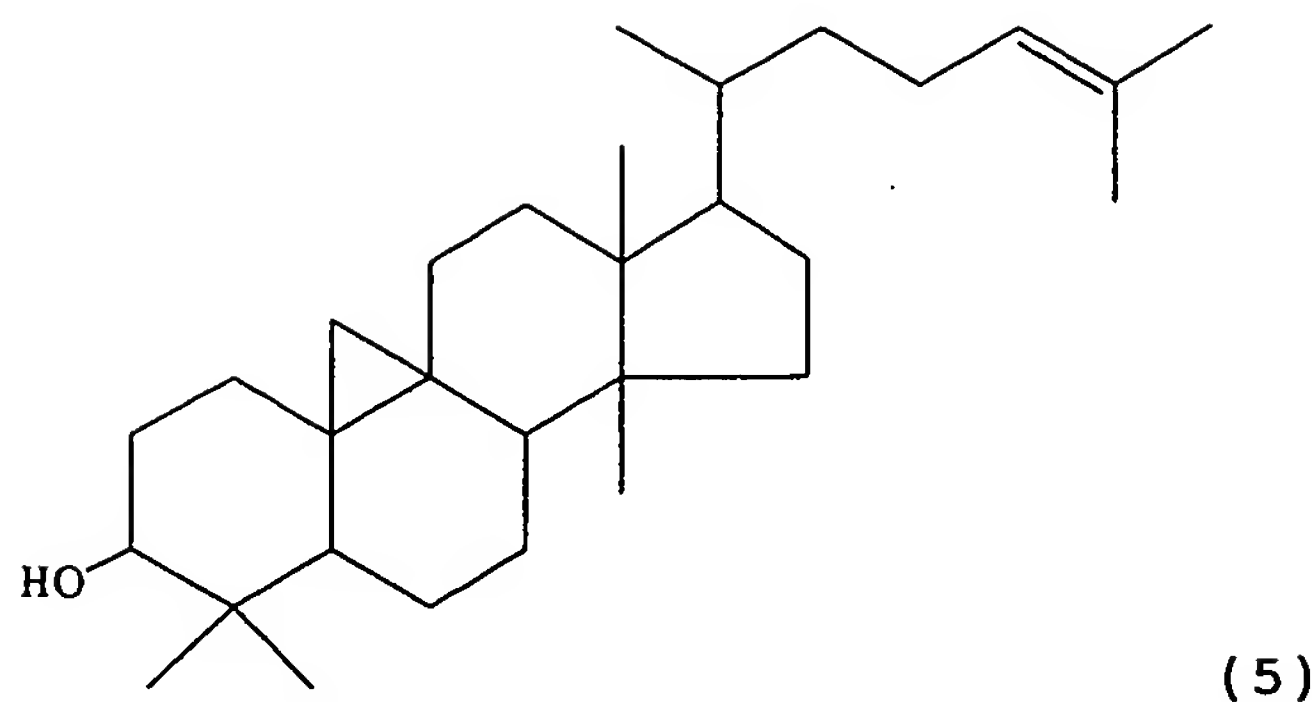


That is, 9,19-cyclolanostan-3-ol is a compound represented by the aforementioned general formula (1) wherein R₂ and R₃ are methyl groups, R₄ is a hydroxyl group, and R₁ is a group represented by the aforementioned formula (i). Further, 24-methylene-9,19-cyclolanostan-3-ol is a compound represented by the aforementioned general formula (1) wherein R₂ and R₃ are methyl groups, R₄ is a hydroxyl group, and R₁ is a group represented by the aforementioned formula (vi).

The compound of the present invention may be cycloartenol (formula (5)) or 24-methylcycloartanol (formula (7)). Both of these compounds are compounds represented by the aforementioned general formula (1)

wherein R2 and R3 are methyl groups, R4 is a hydroxyl group, and R1 is a group represented by the aforementioned formula (vii) in cycloartenol or a group represented by the aforementioned formula (ii) ($R_a = CH_3$, $R_b = H$) in 24-methylcycloartanol.

The compound of the present invention can be chemically produced by a known production method. For example, methods for producing cycloartenol (formula (5)) and 24-methylenecycloartanol (trivial name of 24-methylene-9,19-cyclolanostan-3-ol, formula (4)) have been disclosed in Japanese Patent Laid-open No. 57-018617, and a method for producing cycloartenol ferulate (formula (6)) from γ -oryzanol and a method for synthesizing a compound using a hydrolysate thereof as a starting material have been disclosed in Japanese Patent Laid-open No. 2003-277269. Further, when the R1 moiety of the general formula (1) contains a double bond, various derivative compounds can be produced by using a technique of converting the double bond portion into an aldehyde by ozone decomposition reaction and binding a phosphonate to it, a technique of adding hydrogen to a double bond portion, or a technique of oxidizing the double bond portion with ozone to convert it to an aldehyde or an acid. Further, the production methods are not limited to chemical synthesis methods, and the compounds may be biologically produced by using a microorganism or the like. Alternatively, they may be produced by using enzymes derived from microorganisms.



The drug or food or drink of the present invention may contain one type or two or more arbitrary types of the aforementioned compounds.

It is known that compounds having the cyclolanostane skeleton are contained in plants of the families *Liliaceae*, *Leguminosae*, *Gramineae*, *Solanaceae*, *Musaceae* and so forth (refer to *Phytochemistry*, U.S.A., 1977, Vol. 16, pp.140-141; *Handbook of phytochemical constituents of GRAS herbs and other economic plants*, 1992, U.S.A., CRC Press; Hager's

Handbuch der Pharmazeutischen Praxis, Vols. 2-6, 1969-1979, Germany, Springer-Verlag Berlin). Accordingly, the compounds can be extracted from these plants using a method such as extraction with an organic solvent or extraction with hot water.

In the present invention, although the compound of the present invention may be those purified by the methods described above etc., a composition such as a plant extract or a fraction thereof may also be used so long as it contains an effective amount of the compound.

Specifically, examples of the plant belonging to the family *Liliaceae* include plants belonging to the genus *Aloe* or *Allium*. Examples of the plants of the genus *Aloe* include *Aloe vera* (*Aloe barbadensis* Miller), *Aloe ferox* Miller, *Aloe africana* Miller, *Aloe arborescen* Miller var. *natalensis* Berger, *Aloe spicata* Baker and so forth.

In the production of the compound of the present invention or a composition containing the same, although the whole of the aforementioned plant may be used, it is preferable to use mesophyll (clear gel portion) thereof. Such a plant or a part thereof is disrupted by using a homogenizer or the like and thereby liquefied, and the disruption product is extracted by using an organic solvent or hot water. Examples of the organic solvent include alcohols such as methanol, ethanol and butanol; esters such as methyl acetate, ethyl acetate, propyl acetate and butyl acetate; ketones such as acetone and methyl isobutyl ketone; ethers such as diethyl ether and petroleum ether; hydrocarbons such as hexane, cyclohexane, toluene and benzene; halogenated hydrocarbons such as carbon tetrachloride, dichloromethane and chloroform; heterocyclic compounds such as pyridine; glycols such as ethylene

glycol; polyhydric alcohols such as polyethylene glycol; nitrile solvents such as acetonitrile, mixtures of these solvents and so forth. Further, these solvents may be anhydrous or hydrous. Among these solvents, ethyl acetate/butanol mixture (3:1) and chloroform/methanol mixture (2:1) are particularly preferred.

As the extraction method, a method used for usual extraction of a plant component can be used. Usually used is, for example, a method of refluxing 1 to 300 parts by mass of an organic solvent with 1 part by mass of fresh plant or dried plant with heating at a temperature at or below the boiling point of the solvent and stirring or shaking, or a method of performing extraction by ultrasonication at room temperature. By isolating insoluble matters from the extraction liquor using a suitable method such as filtration or centrifugation, a crude extract can be obtained.

The crude extract can be purified by various types of chromatography such as normal or reverse phase silica gel column chromatography. When a gradient of chloroform/methanol mixture is used in normal phase silica gel column chromatography as an elution solvent, the compound of the present invention is eluted with a mixing ratio of chloroform:methanol = about 25:1. Further, when a hexane/ethyl acetate mixture (4:1) is used in reverse phase silica gel column chromatography as an elution solvent, the compound of the present invention is eluted in a fraction eluted at an early stage.

The obtained fraction can be further purified by HPLC or the like.

Further, the compound used for the present invention may also be produced by a chemical synthesis method or a

biological or enzymatic method using microorganisms, enzymes or the like.

The structure of the compound of the present invention can be confirmed by, for example, mass spectrometry (MS), nuclear magnetic resonance (NMR) spectroscopy or the like.

The compound of the present invention has an action of lowering the hemoglobin Alc level, and as a result, it can control the blood glucose level over a long period of time. Therefore, it can be used as an active ingredient of a drug or food or drink for improving hyperglycemia.

Furthermore, because leaf-skin of Aloe vera contains barbaloin and aloe-emodin having a laxative action, it is conventionally considered to be unfavorable as a drug or food or drink for which laxative action is not expected. Therefore, it is preferred that the composition containing the compound of the present invention does not contain these ingredients. Further, mesophyll of Aloe vera and a disruption product thereof may also be used as an active ingredient of a hyperglycemia improving agent.

The compound of the present invention can be used as an active ingredient of the drug or food or drink of the present invention as it is. Further, an organic solvent extract or a hot water extract of a plant or a fraction thereof containing the compound of the present invention (hereinafter referred to as "extract etc.") may also be used as an active ingredient of the drug or food or drink. In this case, the aforementioned extract etc. to be contained in the drug preferably contains 0.001 to 10% by dry mass, more preferably 0.01 to 1% by dry mass, particularly preferably 0.05 to 1% by dry mass, of the compound of the present invention. Further, the

aforementioned extract etc. to be contained in the food or drink preferably contains 0.0001 to 1% by dry mass, more preferably 0.001 to 1% by dry mass, particularly preferably 0.005 to 1% by dry mass, of the compound of the present invention. The aforementioned extract etc. may contain 2 or more types of the compound of the present invention. Further, the aforementioned extract etc. may be a solution, or can also be lyophilized or spray-dried in a conventional manner and stored or used as powder.

As the drug of the present invention, the compound of the present invention or a composition containing the same such as extract etc. per se, or those combined with a pharmaceutically acceptable carrier can be orally or parenterally administered to a mammal including human. In the drug of the present invention, the compound of the present invention may be a pharmaceutically acceptable salt. Examples of the pharmaceutically acceptable salt include both metal salts (inorganic salts) and organic salts including, for example, those listed in "Remington's Pharmaceutical Sciences," 17th edition, p.1418, 1985. Specific examples thereof include, but not limited to, inorganic acid salts such as hydrochloride, sulfate, phosphate, diphosphate, and hydrobromate, and organic acid salts such as malate, maleate, fumarate, tartarate, succinate, citrate, acetate, lactate, methanesulfonate, p-toluenesulfonate, pamoate, salicylate and stearate. Furthermore, the salt may be a salt with a metal such as sodium, potassium, calcium, magnesium and aluminum or a salt with an amino acid such as lysine. Furthermore, solvates such as hydrates of the aforementioned compound or pharmaceutically acceptable salts thereof also fall within the scope of the present invention.

Dosage form of the drug of the present invention is not particularly limited and can be suitably selected depending on the therapeutic purpose. Specific examples thereof include tablet, pill, powder, solution, suspension, emulsion, granules, capsule, syrup, suppository, injection, ointment, patch, eye drop, nasal drop and so forth. For the preparation, additives generally used in usual hyperglycemia improving drugs as pharmaceutical carriers such as excipients, binders, disintegrating agents, lubricants, stabilizers, flavoring agents, diluents, surfactants and solvents for injection can be used. Further, so long as the effect of the present invention is not degraded, the compound of the present invention, or an extract etc. containing the same can be used in combination with other drugs having a hyperglycemia improving effect.

Although the amount of the compound of the present invention or an extract etc. containing the same contained in the drug of the present invention is not particularly limited and can be suitably selected, the amount in the pharmaceutical preparation may be, for example, 0.001 to 10% by mass, preferably 0.01 to 1% by mass, particularly preferably 0.05 to 1% by mass, in terms of the amount of the compound of the present invention.

The drug of the present invention is useful for a therapeutic or prophylactic treatment of a disease resulted from hyperglycemic conditions such as diabetes and its associated symptoms and conditions (likelihood of developing diabetes or related conditions). In particular, it can also be used to prevent onset of diabetes mellitus from hyperglycemic conditions. Furthermore, the drug of the present invention can cure or prevent various diseases, complications and so forth resulted from hyperglycemic

conditions, and reduce risks of these diseases, complications and so forth.

Examples of such various diseases and complications resulted from hyperglycemic conditions include diabetic retinopathy, diabetic nephropathy, diabetic neuropathy, diabetic gangrene, cerebral apoplexy resulted from diabetes mellitus, myocardial infarction resulted from diabetes mellitus and so forth.

The term "hyperglycemic conditions" refers to conditions that the blood glucose levels are out of the normal ranges, and the normal ranges are generally defined as a fasting blood glucose level of 110 mg/dl or lower, a blood glucose level of 160 mg/dl or lower 1 hour after 75 g glucose load, and a blood glucose level of 120 mg/dl or lower 2 hours after the same glucose load (Nihon Rinsho, No. 806, Vol. 1, pp.28-35, 2002). Furthermore, the drug of the present invention is preferably used for a treatment for a patient with a hemoglobin A1c level higher than normal, for example, a hemoglobin A1c level of 5.8% or higher.

The administration time of the drug of the present invention is not particularly limited and can be suitably selected according to the method for treating an objective disease. Furthermore, the administration route is preferably determined depending on the dosage form, age, sex and other conditions of patients, severity of symptoms of patients and so forth.

The dose of the active ingredient in the drug of the present invention is suitably selected depending on the dosing regimen, age, sex, severity of disease, other conditions of patients and so forth. The amount of the compound of the present invention as an active ingredient is usually selected from the range of, preferably 0.001 to

50 mg/kg/day, more preferably 0.01 to 1 mg/kg/day, as a tentative dose. Further, when an extract etc. containing the compound of the present invention is used, the dry weight of the extract etc. is selected from the range of, preferably 0.1 to 1000 mg/kg/day, more preferably 1 to 100 mg/kg/day, as a tentative amount. In any case, the dose can be ingested, in a day, once or several times as divided portions.

The compound of the present invention or the extract etc. containing the same can be added to food or drink. The form and property of the food or drink are not particularly limited so long as the effect of the active ingredient is not degraded, and the food or drink can be orally ingested, and it can be produced in a conventional manner by using raw materials usually used for food or drink except that the aforementioned active ingredient is added.

The amount of the compound of the present invention or the extract etc. containing the same contained in the food or drink of the present invention is not particularly limited and can be suitably selected. For example, the compound of the present invention or the extract etc. containing the same is contained in food or drink in an amount of 0.0001 to 1% by mass, preferably 0.001 to 1% by mass, particularly preferably 0.005 to 1% by mass, in terms of the amount of the compound of the present invention.

The food or drink of the present invention can be used for various applications utilizing the hyperglycemia improving effect. For example, it can be used for applications as food or drink suitable for those who are getting concerned about their blood glucose levels, food or drink useful for decreasing or eliminating risk factors of

lifestyle-related diseases such as diabetes mellitus, and so forth.

As for the food or drink of the present invention, the expression "improvement of hyperglycemia" means that improvement or prevention of various health damages resulted from hyperglycemia, and "prevention of hyperglycemia," "suppression of increase in blood glucose level," "improvement of increase in blood glucose level," "prevention of increase in blood glucose level," "improvement of high hemoglobin Alc level" and so forth are exemplified in the present invention as terms having a meaning similar to that of the aforementioned "improvement of hyperglycemia".

Furthermore, the food or drink of the present invention is useful for a prophylactic treatment of a disease resulted from hyperglycemic conditions such as diabetes and its associated symptoms and conditions (likelihood of developing diabetes or related conditions). In particular, it can also be used to prevent onset of diabetes from hyperglycemic conditions. Furthermore, the food or drink of the present invention can be used for a prophylactic treatment of various diseases, complications and so forth resulted from hyperglycemic conditions and can decrease risks of these diseases, complications and so forth.

Examples of such various diseases and complications resulted from hyperglycemic conditions include diabetic retinopathy, diabetic nephropathy, diabetic neuropathy, diabetic gangrene, cerebral apoplexy resulted from diabetes mellitus, myocardial infarction resulted from diabetes mellitus and so forth.

The food or drink of the present invention is

preferably marketed as food or drink attached with an indication that the food or drink is used for improving hyperglycemia, for example, "food or drink containing a compound having hyperglycemia improving effect indicated as 'For improving hyperglycemia,'" "food or drink containing a plant extract indicated as 'For improving hyperglycemia,'" "food or drink containing Aloe vera extract indicated as 'For improving hyperglycemia'" and so forth.

Because the compound of the present invention, the composition containing the same and others have a hyperglycemia improving effect, it is considered that the indication of "improvement of hyperglycemia" also means "suppression of increase in blood glucose level." Therefore, the food or drink of the present invention can be indicated as "For suppressing increase in blood glucose level." That is, the aforementioned indication of "For improvement of hyperglycemia" may be an indication of "For suppression of increase in blood glucose level."

The wording used for such an indication as mentioned above is not necessarily be limited to the expression "For improvement of hyperglycemia" or "For suppression of increase in blood glucose level", and any other wording expressing the effect of improving hyperglycemia or suppressing increase in blood glucose level of course falls within the scope of the present invention. As such a wording, for example, an indication based on various uses allowing consumers to recognize the effect of improving hyperglycemia or suppressing increase in blood glucose level is also possible. Examples include, for example, indications of "Suitable for those who are getting concerned with blood glucose levels", "Useful for decrease or elimination of risk factors (risks) of lifestyle-related

diseases such as diabetes mellitus", and so forth.

The aforementioned term "indication" include all actions for informing consumers the aforementioned use, and any indications reminding or analogizing the aforementioned use fall within the scope of the "indication" of the present invention regardless of purpose, content, objective article, medium etc. of the indication. However, the indication is preferably made with an expression that allows consumers to directly recognize the aforementioned use. Specific examples include actions of indicating the aforementioned use on goods or packages of goods relating to the food or drink of the present invention, actions of assigning, delivering, displaying for the purpose of assigning or delivering or importing such goods or packages of goods indicated with the aforementioned use, displaying or distributing advertisements, price lists or business papers relating the goods with indicating the aforementioned use, or providing information including those as contents with indicating the aforementioned use by an electromagnetic method (Internet etc.) and so forth.

The indication is preferably an indication approved by the administration etc. (for example, an indication in a form based on an approval, which is qualified on the basis of any of various legal systems provided by the administration), and it is particularly preferably an indication on advertisement materials at the sales spots such as packages, containers, catalogs, pamphlets and POPs, others documents and so forth.

Examples of the indication further include, for example, indications as health food, functional food, enteric nutritive food, food for special dietary uses, food with nutrient function claims, quasi-drug and so forth as

well as indications approved by the Ministry of Health, Labor and Welfare, for example, indications approved on the basis of the system of food for specified health uses and similar systems. Examples of the latter include indications as food for specified health uses, indications as food for specified health uses with qualified health claims, indications of influence on body structures and functions, indications of reduction of disease risk claims and so forth, and more precisely, typical examples include indications as food for specified health uses (especially indications of use for health) provided in the enforcement regulations of Health Promotion Law (Japan Ministry of Health, Labor and Welfare, Ministerial ordinance No. 86, April 30, 2003) and similar indications.

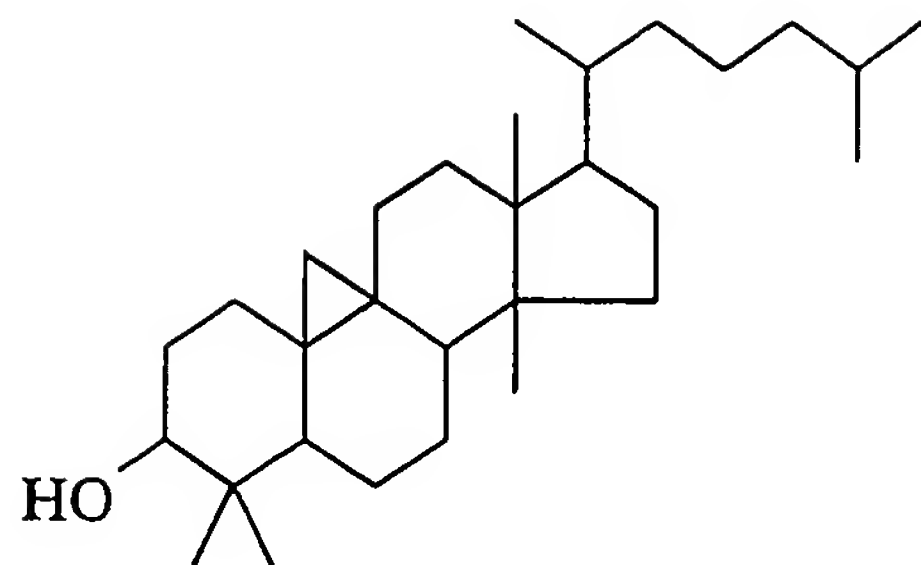
EXAMPLES

The present invention will be explained more specifically with reference to the following examples. However, the scope of the present invention is not limited to the following examples.

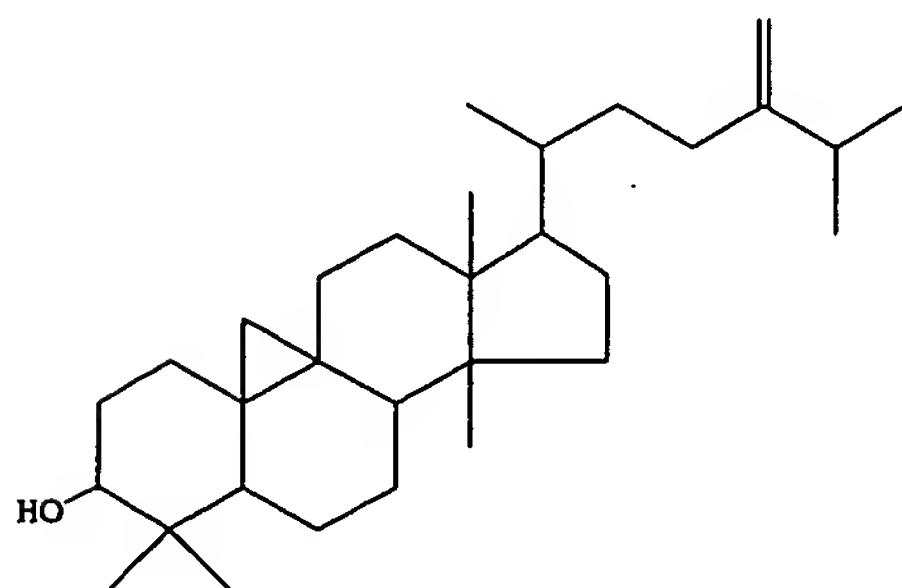
Preparation examples of compounds having the lanostane skeleton will be mentioned below.

[Preparation Example 1]

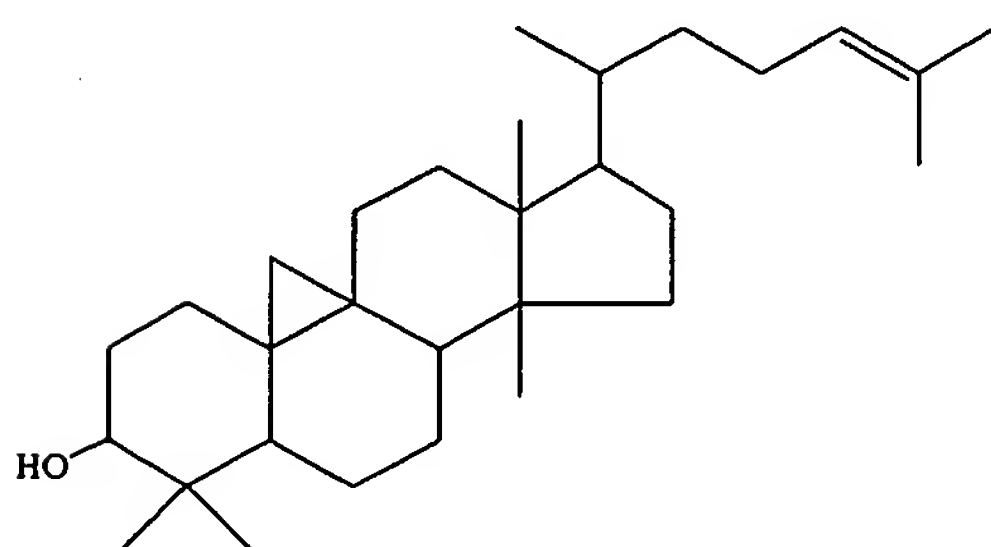
9,19-Cyclolanostan-3-ol(formula (3)), 24-methylene-9,19-cyclolanostan-3-ol (formula (4)), cycloartenol (formula (5)) and 24-methylcycloartanol (formula (7)) were prepared by the method described below.



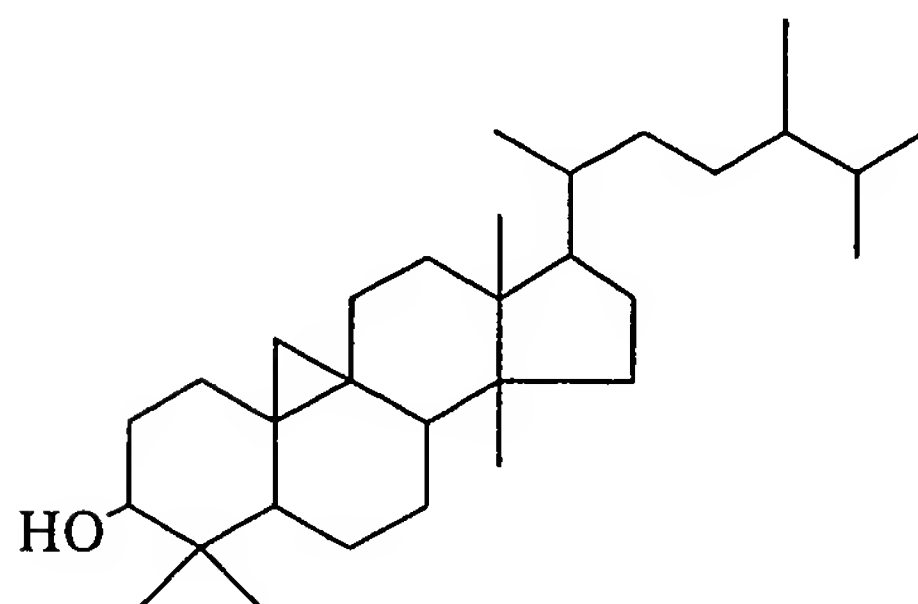
(3)



(4)



(5)



(7)

To 8.0 g of γ -oryzanol (Oryza Oil & Chemical Co., Ltd.) was added 250 ml of distilled water, 50 g of sodium

hydroxide, 150 ml of isopropanol, 150 ml of ethanol and 150 ml of methanol, and the mixture was refluxed with heating for 2 hours by using a mantle heater. After the reaction, the reaction mixture was poured into 1300 ml of water, and the produced white precipitates were isolated by suction filtration. To wash off the remaining alkali, the residue obtained by the filtration was suspended in 1000 ml of water, and then collected by suction filtration again. This procedure was repeated twice, and the finally obtained residue was lyophilized under reduced pressure to obtain 5.91 g of an oryzanol hydrolysate. This hydrolysate was purified by HPLC to obtain 2435 mg of cycloartenol and 1543 mg of 24-methylene-9,19-cyclolanostan-3-ol.

The obtained cycloartenol was used to synthesize 9,19-cyclolanostan-3-ol. In an amount of 302 mg of cycloartenol, 150 ml of isopropanol and 1.0 g of powdery 5% palladium/carbon catalyst were charged into a sealed autoclave, the internal atmosphere was replaced with a nitrogen gas, and then a hydrogen gas was introduced with applying 3 kg/cm² of pressure. The mixture was heated with stirring, and when the temperature reached 50°C, the hydrogen pressure was adjusted to 5 kg/cm². With supplementing hydrogen for the absorbed hydrogen to maintain the pressure, the reaction was allowed for 6 hours. The reaction mixture was filtered to remove the catalyst, concentrated and then purified by silica gel column chromatography (developing solvent: 100% chloroform) to obtain 275 mg of 9,19-cyclolanostan-3-ol. 24-Methylcycloartanol was synthesized by using 24-methylene-9,19-cyclolanostan-3-ol as a starting material. In an amount of 78 mg of 24-methylene-9,19-cyclolanostan-3-ol, 150 ml of isopropanol and 1.0 g of powdery 5%

palladium/carbon catalyst were charged into a sealed autoclave, the internal atmosphere was replaced with a nitrogen gas, and then a hydrogen gas was introduced with applying 3 kg/cm² of pressure. Then, the mixture was heated with stirring, and when the temperature reached 50°C, the hydrogen pressure was adjusted to 5 kg/cm². With supplementing hydrogen for the absorbed hydrogen to maintain the pressure of 5 kg/cm², the reaction was allowed for 6 hours. The reaction mixture was filtered to remove the catalyst, concentrated and then purified by silica gel column chromatography (developing solvent: 100% chloroform) to obtain 69 mg of 24-methylcycloartanol.

Preparation examples of extracted compositions containing a compound having the cyclolanostane skeleton using Aloe vera (*Aloe barbadensis* Miller) as a starting material will be described below.

[Preparation Example 2]

In an amount of 100 kg of hulled Aloe vera (*Aloe barbadensis* Miller) was liquefied by using a homogenizer, added with 100 L of an ethyl acetate ester/butanol mixture (3:1) and stirred. The mixture was left overnight to separate the ethyl acetate ester/butanol mixture and the aqueous layer, and the ethyl acetate ester/butanol mixture was recovered. The extracted composition containing a compound having the cyclolanostane skeleton, which was obtained by concentrating the ethyl acetate ester/butanol mixture under reduced pressure, weighed 13.5 g. LC-MS measurement of this composition revealed that the content of 9,19-cyclolanostan-3-ol was 10 mg, and the content of 24-methylene-9,19-cyclolanostan-3-ol was 70 mg.

[Preparation Example 3]

In an amount of 1 kg of Aloe vera powder was added with 10 L of a chloroform/methanol mixture (2:1) and immersed overnight in the mixture at room temperature, and then the chloroform/methanol mixture was recovered. The organic solvents were completely removed from this mixture at 28°C to obtain 83 g of a composition containing a compound having the cyclolanostane skeleton. LC-MS measurement of this composition revealed that the content of 9,19-cyclolanostan-3-ol was 25.8 mg, and the content of 24-methylene-9,19-cyclolanostan-3-ol was 24 mg.

[Test Example 1]

This test was performed in order to determine the hemoglobin Alc lowering action of a compound having the cyclolanostane skeleton.

(1) Preparation of sample

The 9,19-cyclolanostan-3-ol and 24-methylene-9,19-cyclolanostan-3-ol produced in Preparation Example 1 mentioned above were used as test samples 1 and 2, respectively.

(2) Test method

As type-II diabetes model mice, 6-week old male db/db mice (purchased from Clea Japan, Inc.) were used in this test. These mice were divided into groups, each consisting of 7 animals. Each test sample was dissolved in DMSO, and the concentration was adjusted to 1 µg/mL with physiological saline. The final DMSO concentration was adjusted to 0.2%. A solution that did not contain either of the test samples was used as a negative sample solution. The type-II diabetes model mice were administered with 1 mL

of either of the test sample solutions or the negative sample solution once a day everyday with a sonde for 45 consecutive days. On the 43rd day of the repetitive administration, hemoglobin A1c levels were measured by using DCA 2000 (Bayer-Sankyo Co., Ltd.).

(3) Test results (hemoglobin A1c levels)

The hemoglobin A1c levels on the 43rd day of the repetitive administration of the sample solutions are shown in Table 1. In comparison with the hemoglobin A1c levels after the administration of the negative sample, lowering effects of 8.2 and 14.5% were observed after the administration of the test samples 1 and 2, respectively. Further, there was no case showing acute hypoglycemic conditions after the administration during the administration period, and no adverse side effect symptom was observed from viewpoints of body weight and pathological findings.

Table 1

Samples	Hemoglobin A1c levels (%) 43rd day from the administration	p value against negative sample
Test sample 1	91.8±19.6	<0.3627>
Test sample 2	85.6±9.3	<0.0073>
negative sample	100	

[Test Example 2]

This test was performed in order to compare the hemoglobin A1c lowering action of the compounds of the present invention with that of an antidiabetic drug used in the clinical practice.

(1) Preparation of sample

The same test sample 1 (9,19-cyclolanostan-3-ol) and test sample 2 (24-methylene-9,19-cyclolanostan-3-ol) used in Test Example 1 were used.

(2) Test method

As type-II diabetes model mice, 6-week old male db/db mice (purchased from Clea Japan, Inc.) were used. These mice were divided into groups, each consisting of 7 animals. Each test sample was dissolved in DMSO, and the concentration was adjusted to 1 $\mu\text{g/mL}$ with physiological saline. The final DMSO concentration was adjusted to 0.2%. A solution that did not contain any of the test samples was used as a negative sample. Further, as a control sample 1, ACTOS[®] tablets (Takeda Pharmaceutical Co., Ltd.) were ground in a mortar and dissolved in physiological saline so that the concentration of pioglitazone hydrochloride as the active ingredient should become 7.5 $\mu\text{g/mL}$. The type-II diabetes model mice were orally administered with 1 mL of each test sample solution, control sample 1 solution or negative sample solution once a day everyday with a sonde for 22 consecutive days. On the 23rd day from the start of the administration, hemoglobin A1c levels were measured by using DCA 2000 (Bayer-Sankyo Co., Ltd.).

(3) Test results (hemoglobin A1c levels)

The measurement results of hemoglobin A1c levels on the 23rd day from the start of the administration are shown in Table 2. In comparison with the hemoglobin A1c levels after the administration of the negative sample, statistically significant decreases of 15 to 18% in the

hemoglobin Alc levels were observed after the administration of either the test sample 1 or 2. In contrast, decreases of only about 0.8% were observed after the administration of the control sample 1, and no statistically significant effect was obtained either. Further, there was no case showing acute hypoglycemic conditions during the administration period or after the administration, and no adverse side effect symptom was observed from viewpoints of body weight and pathological findings.

Table 2

Samples	Hemoglobin Alc levels (%) 23rd day from the administration	p value against negative sample
Test sample 1	85.4 ± 11.2	<0.009>
Test sample 2	82.0 ± 7.9	<0.003>
Control sample 1	92.1 ± 0.2	<0.2>
negative sample	100	

[Test Example 3]

In this test, examined was the effective concentration of 24-methylene-9,19-cyclolanostan-3-ol, which showed the most potent hemoglobin Alc lowering action in Test Example 1.

(1) Preparation of sample

The 24-methylene-9,19-cyclolanostan-3-ol produced in Preparation Example 1 mentioned above was used as a test sample 3.

(2) Test method

In this test, 6-week old male db/db mice (purchased

from Clea Japan, Inc.) were used as type-II diabetes model mice. These mice were divided into groups, each consisting of 7 animals. The test sample 3 was dissolved in DMSO, and the concentration of 24-methylene-9,19-cyclolanostan-3-ol was adjusted to 0.1 or 1 µg/ml with physiological saline. The final DMSO concentration was adjusted to 0.2%. A solution that did not contain the test sample was used as a negative sample. The type-II diabetes model mice were administered with 1 mL of the test sample 3 solution of either concentration or negative sample solution once a day everyday with a sonde for 40 consecutive days. On the 41st day from the start of the repetitive administration, hemoglobin Alc levels were measured by using DCA 2000 (Bayer-Sankyo Co., Ltd.).

(3) Test results (hemoglobin Alc levels)

The hemoglobin Alc levels on the 41st day from the start of the repetitive administration of the sample solutions are shown in Table 3. In comparison with the hemoglobin Alc levels observed after the administration of the negative sample, a lowering effect of about 1.9% for the hemoglobin Alc levels was observed after the administration of the test sample 3 at a dose of 0.1 µg/day, and a significant lowering effect of 18.4% was observed after the administration at a dose of 1 µg/day.

Table 3

Samples	Hemoglobin Alc levels (%) 41st day from the administration	p value against negative sample
Test sample 3 0.1 μ g	98.1 \pm 22.3	<0.8387>
Test sample 3 1 μ g	81.6 \pm 14.6	<0.0235>
Negative sample	100	

[Test Example 4]

This test was performed in order to evaluate the hemoglobin Alc lowering action and hyperglycemia improving effect activity of compounds having the cyclolanostane skeleton.

(1) Preparation of samples

The 9,19-cyclolanostan-3-ol and 24-methylene-9,19-cyclolanostan-3-ol produced in Preparation Example 1 mentioned above were used as test samples 1 and 2, respectively. Further, β -sitosterol (Tama Biochemical Co., Ltd.) was used as a control sample.

(2) Test method

In this test, 6-week old male db/db mice (purchased from Clea Japan, Inc.) were used as type-II diabetes model mice. These mice were divided into groups, each consisting of 7 animals. Each sample was dissolved in DMSO, and the concentrations of 9,19-cyclolanostan-3-ol and 9,19-cyclolanost-25-en-3-ol were adjusted to 1 μ g/mL with physiological saline to obtain test sample 1 solution and test sample 2 solution. Further, concentration of β -sitosterol was adjusted to 15 μ g/ml in a similar manner to

obtain a control sample solution. The final DMSO concentration was adjusted to 0.2%. A solution that did not contain either the test samples was used as a negative sample. The type-II diabetes model mice were administered with 1 mL of one of the test sample solutions, control sample solution and negative sample solution once a day everyday with a sonde for 40 consecutive days. Fasting blood glucose levels and random blood glucose levels were measured over time by using Antsense II (Bayer-Sankyo Co., Ltd.). The fasting blood glucose levels were measured after 15 hours of fasting.

(3) Test results

Changes over time in random blood glucose levels and fasting blood glucose levels during the repetitive administration period of each sample are shown in Figs. 1 and 2. Whereas rapid increase were observed in both the random blood glucose levels and fasting blood glucose levels in the mice administered with the negative sample or control sample, effect of suppressing the increases in blood glucose levels was clearly observed in the mice repeatedly administered with the test sample 1 or 2.

Further, there was no case showing acute hypoglycemic conditions after the administration during the administration period, and no adverse side effect symptom was observed from viewpoints of body weight and pathological findings.

[Test Example 5]

This test was performed in order to evaluate the hemoglobin Alc lowering action of an extracted composition derived from Aloe vera (*Aloe barbadensis* Miller) and

containing a compound having the cyclolanostane skeleton.

(1) Preparation of sample

The extracted composition containing a compound having the cyclolanostane skeleton prepared in Preparation Example 2 mentioned above was used as a test sample 4.

(2) Test method

In this test, 6-week old male db/db mice (purchased from Clea Japan, Inc.) were used as type-II diabetes model mice. These mice were divided into groups, each consisting of 7 animals. The test sample was dissolved in DMSO, and the solid content of the extracted composition was adjusted to 25 or 250 $\mu\text{g/ml}$ with physiological saline. The final DMSO concentration was adjusted to 0.2%. A solution that did not contain the test sample was used as a negative sample solution. The type-II diabetes model mice were administered with 1 mL of test sample 4 solution at either concentration or negative sample solution once a day everyday with a sonde for 34 consecutive days. On the 35th day from the start of the administration, hemoglobin A1c levels were measured by using DCA 2000 (Bayer-Sankyo Co., Ltd.).

(3) Test results (hemoglobin A1c levels)

The hemoglobin A1c levels on the 35th day from the start of the repetitive administration are shown in Table 4. In comparison with the hemoglobin A1c levels observed after the administration of the negative sample, decreases in hemoglobin A1c levels were observed after the administration of the test sample 4 solution of either solid content of 25 or 250 $\mu\text{g/ml}$ in the extracted

composition, indicating existence of statistically significant blood glucose level control effect over a long period of time. Further, during the administration period and after the administration, there was no case showing adverse side reaction symptoms or acute hypoglycemic conditions, and no problem concerning safety was observed from viewpoints of body weight and pathological findings.

Table 4

Samples	Hemoglobin Alc levels (%) 35rd day from the administration	p value against negative sample
Test sample 4 25 μ g	82.6 ± 7.1	$<0.001>$
Test sample 4 250 μ g	84.9 ± 8.2	$<0.007>$
Negative sample	100	

Industrial Applicability

The drug and food or drink of the present invention can be safely administered or ingested without causing acute hypoglycemia and have a long-term blood glucose level control action lowering the hemoglobin Alc level. Further, the active ingredient of the drug and food or drink of the present invention can be produced from a plant that can be safely ingested from an experiential viewpoint for food and is readily available, for example, a plant of the family *Liliaceae* such as *Aloe vera* (*Aloe barbadensis* Miller).